

3. What changes did we make to GRS 2.7?

We added two items to cover records of vaccine attestations for Federal employees and contractors (item 063) and visitors (item 064), and two items to cover symptom screening and testing records for Federal employees (item 065) and contractors and visitors (item 66).

4. What changes did we make to GRS 4.2?

We reduced the retention period of item 100. It previously directed agencies to retain the records for 30 years after completing a declassification review. Now, agencies may destroy records documenting a declassification review immediately upon either of two subsequent events: The agency conducts another declassification review or the agency transfers the reviewed records to NARA. We altered the retention period in response to a request from the Department of State, which pointed out that the previous instruction could result in agencies being required to retain records documenting the declassification process until as late as 105 years after the records were created.

5. What changes did we make to GRS 4.4?

We modified the background information to clarify that the schedule applies to library and information centers within agencies, but not to stand-alone libraries, such as the Library of Congress, or national libraries.

6. What changes did we make to GRS 5.3?

In the first sentence of the Background Information, we changed the generic word “sensitive” to a term with a precise definition: “controlled unclassified.”

7. What changes did we make to GRS 5.6?

We updated this schedule to further clarify that it does not include records related to Federal law enforcement and Federal correctional activities and that this exclusion includes border and transportation security and immigration and naturalization services. We changed the schedule title “Security Management Records” to help with this distinction.

We altered item 010 to clarify the subject matter as security *management* and expanded the description’s list of examples. We removed the bullet for standard operating procedures manuals, as they are properly covered by GRS 5.7, item 030.

We revised items 030, 090, and 100 to clarify that they do not cover records related to Federal law enforcement and correctional activities including border and transportation security and immigration and naturalization services. Item 090 was revised to make it clear that it does not apply to videos of accidents or incidents or video surveillance of accidents or incidents in Federal facilities or facilities operated by contractors on behalf of the Federal Government.

We revised item 120’s disposition instruction to be more concise. We revised item 130’s title to clarify that it covers all manner of temporary access identification records. We changed the term “sensitive data” to “controlled unclassified information” in items 180 and 181.

8. What changes did we make to GRS 5.7?

We revised this schedule to clarify that it applies only to records related to management and oversight of agency administrative functions. This included changing the name of the schedule to “Administrative Management and Oversight Records” and updating the background information to clarify that it applies only to management of administrative functions, not functions related to agency mission. The new background section specifically excludes records related to agency strategic planning and performance management.

We altered item 010’s title and revised the list of examples to remove generic records types that are arguably not “management controls.”

We revised the title of item 040 and added a sentence to the description to clarify that the item applies only to requirements for reports related to administrative activities. We also added an exclusion to clarify that item 040 does not cover the reports themselves.

We eliminated from item 050’s list of included records reports that are not specific to administrative activities, such as Performance and Accountability Reports (PAR). We also added an exclusion to make it clear that mandatory reports related to non-administrative matters are not covered by this item and must be scheduled by the agency.

9. How do agencies cite GRS items?

When you send records to an FRC for storage, you should cite the records’ legal authority—the “DAA” number—in the “Disposition Authority” column of the table. Please also include schedule and item number. For example, “DAA–

GRS–2017–0007–0008 (GRS 2.2, item 070).”

10. Do agencies have to take any action to implement these GRS changes?

NARA regulations (36 CFR 1226.12(a)) require agencies to disseminate GRS changes within six months of receipt.

Per 36 CFR 1227.12(a)(1), you must follow GRS dispositions that state they must be followed without exception.

Per 36 CFR 1227.12(a)(3), if you have an existing schedule that differs from a new GRS item that does *not* require being followed without exception, and you wish to continue using your agency-specific authority rather than the GRS authority, you must notify NARA within 120 days of the date of this transmittal.

If you do not have an already existing agency-specific authority but wish to apply a retention period that differs from that specified in the GRS, you must submit a records schedule to NARA for approval via the Electronic Records Archives.

David S. Ferriero,

Archivist of the United States.

[FR Doc. 2022–07041 Filed 4–1–22; 8:45 am]

BILLING CODE 7515–01–P

NATIONAL SCIENCE FOUNDATION

Sunshine Act Meetings

The National Science Board’s Executive Committee hereby gives notice of the scheduling of a teleconference for the transaction of National Science Board business, pursuant to the National Science Foundation Act and the Government in the Sunshine Act.

TIME AND DATE: Tuesday, April 5, 2022, from 2:00–3:00 p.m. EDT.

PLACE: This meeting will be held by teleconference through the National Science Foundation.

STATUS: Open.

MATTERS TO BE CONSIDERED: Committee Chair’s opening remarks; approval of Executive Committee minutes of January 26, 2022; and discuss issues and topics for an agenda of the NSB meeting scheduled for May 5–6, 2022.

CONTACT PERSON FOR MORE INFORMATION: Point of contact for this meeting is: Nirmala Kannankutty, 703/292–8000. Members of the public can observe this meeting through a YouTube livestream. The link is <https://youtu.be/3qY4Jf5RAvY>. Information about meeting updates is available from the

NSB website at <https://www.nsf.gov/nsb/meetings/index.jsp#up>.

Chris Blair,

Executive Assistant to the National Science Board Office.

[FR Doc. 2022-07148 Filed 3-31-22; 4:15 pm]

BILLING CODE 7555-01-P

NUCLEAR REGULATORY COMMISSION

[NRC-2021-0218]

Information Collection: Specific Domestic Licenses To Manufacture or Transfer Certain Items Containing Byproduct Material

AGENCY: Nuclear Regulatory Commission.

ACTION: Renewal of existing information collection; request for comment.

SUMMARY: The U.S. Nuclear Regulatory Commission (NRC) invites public comment on the renewal of Office of Management and Budget (OMB) approval for an existing collection of information. The information collection is entitled, "Specific Domestic Licenses to Manufacture or Transfer Certain Items Containing Byproduct Material."

DATES: Submit comments by June 3, 2022. Comments received after this date will be considered if it is practical to do so, but the Commission is able to ensure consideration only for comments received on or before this date.

ADDRESSES: You may submit comments by any of the following methods; however, the NRC encourages electronic comment submission through the Federal rulemaking website:

- *Federal Rulemaking Website:* Go to <https://www.regulations.gov> and search for Docket ID NRC-2021-0218. Address questions about Docket IDs in *Regulations.gov* to Stacy Schumann; telephone: 301-415-0624; email: Stacy.Schumann@nrc.gov. For technical questions, contact the individual listed in the **FOR FURTHER INFORMATION CONTACT** section of this document.

- *Mail comments to:* David Cullison, Office of the Chief Information Officer, Mail Stop: T-6 A10M, U.S. Nuclear Regulatory Commission, Washington, DC 20555-0001.

For additional direction on obtaining information and submitting comments, see "Obtaining Information and Submitting Comments" in the **SUPPLEMENTARY INFORMATION** section of this document.

FOR FURTHER INFORMATION CONTACT: David C. Cullison, Office of the Chief Information Officer, U.S. Nuclear

Regulatory Commission, Washington, DC 20555-0001; telephone: 301-415-2084; email: Infocollects.Resource@nrc.gov.

SUPPLEMENTARY INFORMATION:

I. Obtaining Information and Submitting Comments

A. Obtaining Information

Please refer to Docket ID NRC-2021-0218 when contacting the NRC about the availability of information for this action. You may obtain publicly available information related to this action by any of the following methods:

- *Federal Rulemaking Website:* Go to <https://www.regulations.gov> and search for Docket ID NRC-2021-0218.
- *NRC's Agencywide Documents Access and Management System (ADAMS):* You may obtain publicly available documents online in the ADAMS Public Documents collection at <https://www.nrc.gov/reading-rm/adams.html>. To begin the search, select "Begin Web-based ADAMS Search." For problems with ADAMS, please contact the NRC's Public Document Room (PDR) reference staff at 1-800-397-4209, 301-415-4737, or by email to PDR.Resource@nrc.gov. The supporting statement and NRC Form 653, 653A, 653B are available in ADAMS under Accession Nos. ML22028A014 and ML22028A015.

- *NRC's PDR:* You may examine and purchase copies of public documents, by appointment, at the NRC's PDR, Room P1 B35, One White Flint North, 11555 Rockville Pike, Rockville, Maryland 20852. To make an appointment to visit the PDR, please send an email to PDR.Resource@nrc.gov or call 1-800-397-4209 or 301-415-4737, between 8:00 a.m. and 4:00 p.m. (ET), Monday through Friday, except Federal holidays.

- *NRC's Clearance Officer:* A copy of the collection of information and related instructions may be obtained without charge by contacting the NRC's Clearance Officer, David Cullison, Office of the Chief Information Officer, U.S. Nuclear Regulatory Commission, Washington, DC 20555-0001; telephone: 301-415-2084; email: Infocollects.Resource@nrc.gov.

B. Submitting Comments

The NRC encourages electronic comment submission through the Federal rulemaking website (<https://www.regulations.gov>). Please include Docket ID NRC-2021-0218 in your comment submission.

The NRC cautions you not to include identifying or contact information in comment submissions that you do not

want to be publicly disclosed in your comment submission. All comment submissions are posted at <https://www.regulations.gov> and entered into ADAMS. Comment submissions are not routinely edited to remove identifying or contact information.

If you are requesting or aggregating comments from other persons for submission to the OMB, then you should inform those persons not to include identifying or contact information that they do not want to be publicly disclosed in their comment submission. Your request should state that comment submissions are not routinely edited to remove such information before making the comment submissions available to the public or entering the comment into ADAMS.

II. Background

In accordance with the Paperwork Reduction Act of 1995 (44 U.S.C. Chapter 35), the NRC is requesting public comment on its intention to request the OMB's approval for the information collection summarized below.

1. *The title of the information collection:* Part 32 of title 10 of the *Code of Federal Regulations* (10 CFR), "Specific Domestic Licenses to Manufacture or Transfer Certain Items Containing Byproduct Material."

2. *OMB approval number:* 3150-0001.

3. *Type of submission:* Extension.

4. *The form number, if applicable:* NRC Form 653, 653A, 653B.

5. *How often the collection is required or requested:* There is a one-time submittal of information to receive a certificate of registration for a sealed source and/or device. Certificates of registration for sealed sources and/or devices can be amended at any time. In addition, licensee recordkeeping must be performed on an on-going basis, and reporting of transfer of byproduct material must be reported every calendar year, and in some cases, every calendar quarter.

6. *Who will be required or asked to respond:* All specific licensees who manufacture or initially transfer items containing byproduct material for sale or distribution to general licensees, or persons exempt from licensing, medical use product distributors to specific licensees, and those requesting a certificate of registration for a sealed source and/or device.

7. *The estimated number of annual responses:* 3,038 [2,285 reporting + 349 recordkeepers + 404 third-party recordkeepers].

8. *The estimated number of annual respondents:* 662 (156 NRC licenses, registration certificate holder, and 506